## **Patent Claims**

- 1. A membrane for removing proteases from liquid, comprising a microporous membrane body, characterized by the fact that inhibitors binding selectively with proteases are coupled with chemically activated groups to the membrane bodies (7, 8, 9, 10).
- 2. The membrane according to claim 1, characterized by the fact that an inhibitor biding with acidic proteases is coupled to the membrane body (7).
- 3. The membrane according to claim 2, characterized by the fact that pepstatin is coupled to the membrane body (7).
- 4. The membrane according to claim 1 to 3, characterized by the fact that an inhibitor binding with metalloproteases is coupled to the membrane body (8).
- 5. The membrane according to claim 4, characterized by the fact that bestatin, diprotin or EDTA is coupled to the membrane body (8).
- 6. The membrane according to claim 1 through 5, characterized by the fact that an inhibitor binding with cysteine proteases is coupled to the membrane body (9).
- 7. The membrane according to claim 6, characterized by the fact that antipain, chymostatin, leupeptin or E64 is coupled to the membrane body (9).

- 8. The membrane according to claim 1 through 7, characterized by the fact that an inhibitor binding with serine proteases is coupled to the membrane body (10).
- 9. The membrane according to claim 8, characterized by the fact that TLCK or p-aminobenzamidine is coupled to the membrane body (10).
- 10. Device for removing proteases from biological liquids and pharmaceutical solutions with a plurality of connected membranes, characterized by the fact that the membranes (3, 4, 5, 6) are constructed according to one of the claims 1 through 9.
- 11. Device according to claim 10, characterized by the fact that each of the individual membranes (3, 4, 5, 6) is provided with a membrane body (7, 8, 9, 10) having another coupled inhibitor.

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- 12. The device according to the claim 10 or 11, characterized by the fact that the membranes (3, 4, 5, 6) are built into a suitable housing (2) enabling a sequential flow through the membranes (3, 4, 5, 6).
- 13. A method for removing proteases from biological liquids and pharmaceutical solutions with microfiltration using microporous activated membranes, characterized by the fact that the inhibitors are coupled with chemically activated groups to the membranes (3, 4, 5, 6), wherein the proteases are removed with selective binding.

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